

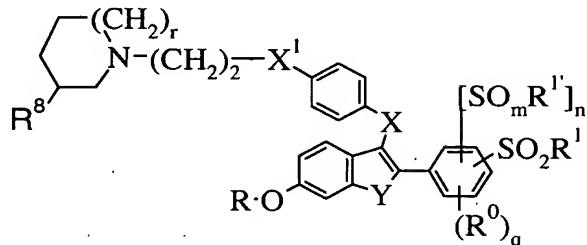
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DT01 Rec'd PCT/PTC 18 JAN 2005

Amendments to the Claims

Without prejudice or disclaimer, this listing of claims will replace all prior versions and listing of claims in this application.

1. (Presently amended) A compound of formula I:



I;

wherein:

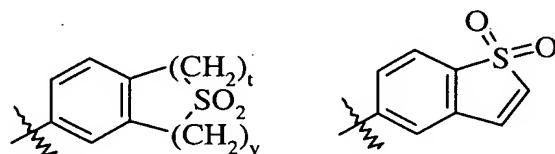
m, q and r are independently 0, 1 or 2;

n is 0 or 1;

R is H or COR²;

R⁰ is independently at each occurrence OH, CF₃, halo, C₁-C₆ alkyl or C₁-C₆ alkoxy;

R¹ and R^{1'} are independently C₁-C₆ alkyl, C₁-C₆ alkoxy, NR³R^{3a}, CF₃ or CH₂CF₃; or when n and q are 0, the -SO₂R¹ moiety may combine with the phenyl ring to which it is attached to form a moiety of formula (a) or (b):



(a)

(b);

wherein t and v are 0, 1 or 2 provided that the sum of t + v must be 2;

R² is C₁-C₆ alkyl; C₁-C₆ alkoxy; NR⁴R⁴; phenoxy; or phenyl optionally substituted with halo;

R³ is C₁-C₆ alkyl or phenyl;

R^{3a} and R⁴ are independently at each occurrence H, C₁-C₆ alkyl, or phenyl;

X is O, CH₂ or CO;

X¹ is O or NR⁵;

R⁵ is H or C₁-C₆ alkyl; and

R⁸ is H or methyl provided that if r is 1 or 2, then R⁸ must be H and that if r is 0, then R⁸ must be methyl; and

Y is S; CH₂CH₂ or CH=CH; or a pharmaceutical acid addition salt thereof.

2. (Original) The compound of claim 1 wherein m is 2; and r is 1 or 2; or a pharmaceutical acid addition salt thereof.

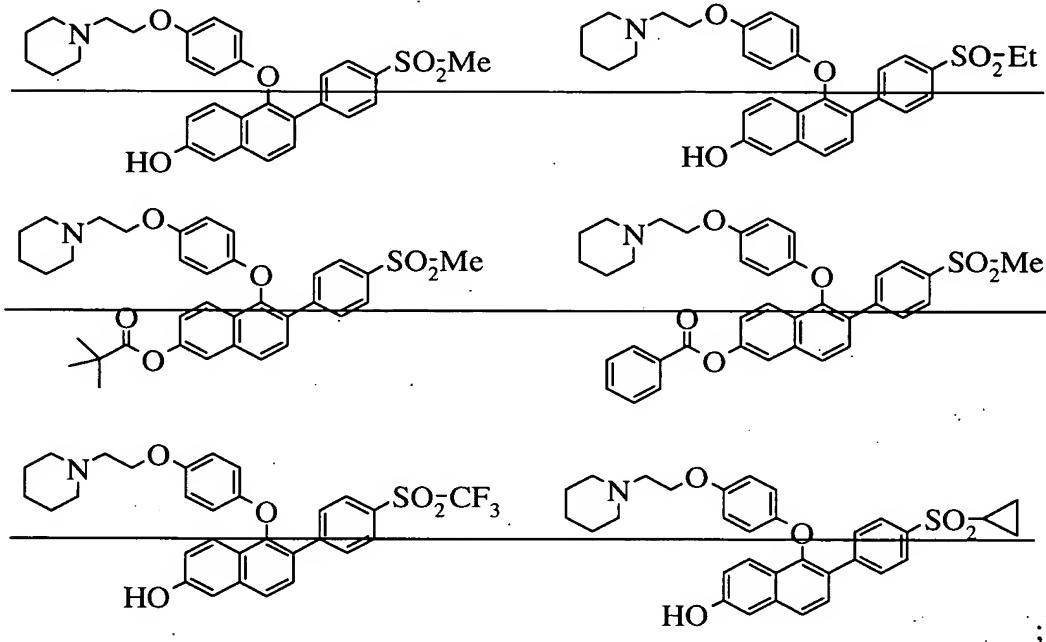
3. (Presently amended) The compound of claim 1 or 2 wherein R² is C₁-C₆ alkyl, NHCH₃ or phenyl and the -SO₂R¹ moiety does not combine with the phenyl ring to which it is attached to form a moiety of formula (a) or (b); or a pharmaceutical acid addition salt thereof.

4. (Presently amended) The compound of ~~any one of claims 1-3~~ claim 3 wherein n is 0; q is 0 or 1; the -SO₂R¹ moiety is at the para-position of the phenyl ring to which it is attached; R⁰ is OH, CF₃, fluoro, chloro, methyl or ethyl; R¹ is methyl, ethyl, n-propyl, isopropyl, cyclopropyl, n-butyl, isobutyl, sec-butyl, t-butyl, cyclobutyl or CF₃; R² is C₁-C₆ alkyl or phenyl; and Y is S or CH=CH; or a pharmaceutical acid addition salt thereof.

5. (Presently amended) The compound of ~~any one of claims 1-4~~ claim 4 wherein X and X¹ are O; or a pharmaceutical acid addition salt thereof.

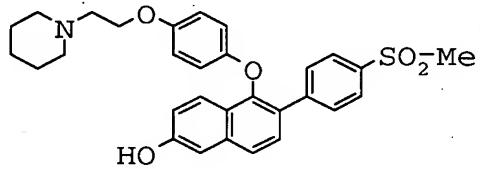
6. (Presently amended) The compound of ~~any one of claims 1-5~~ claim 5 wherein q is 0; R¹ is methyl, ethyl, cyclopropyl or CF₃; and Y is CH=CH; or a pharmaceutical acid addition salt thereof.

7. (Presently amended) The compound of ~~any one of claims 1-6~~ claim 6 wherein R is H selected from the group consisting of:



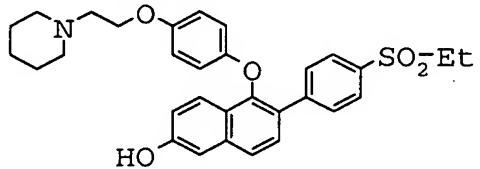
or a pharmaceutical acid addition salt thereof.

8. (Presently amended) The compound of claim 7 which is:



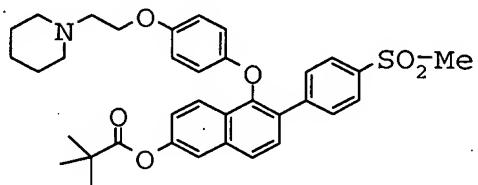
or a pharmaceutical acid addition salt thereof.

9. (Presently amended) The compound of claim 7 which is:



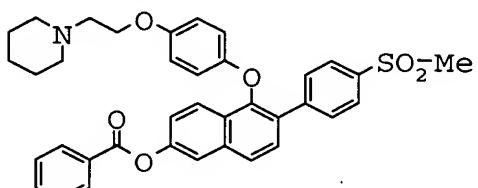
or a pharmaceutical acid addition salt thereof.

10. (Presently amended) The compound of claim 6 which is:



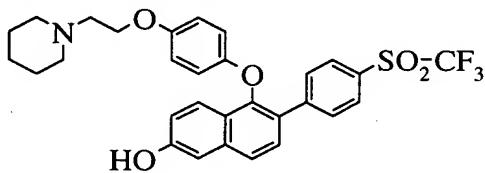
or a pharmaceutical acid addition salt thereof.

11. (Presently amended) The compound of claim 6 which is:



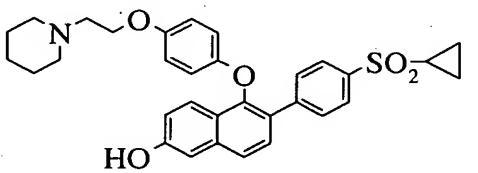
or a pharmaceutical acid addition salt thereof.

12. (Presently amended) The compound of claim 7 which is:



or a pharmaceutical acid addition salt thereof.

13. (Presently amended) The compound of claim 7 which is:



or a pharmaceutical acid addition salt thereof.

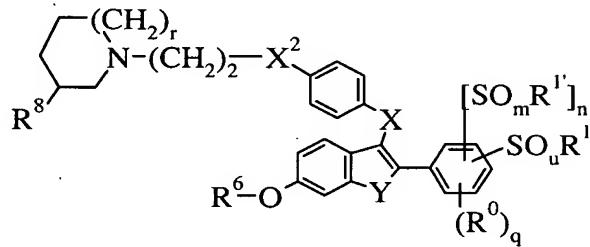
14. (Presently Amended) The compound of ~~any one of claims 1-13~~ claim 8 which is the hydrochloride salt.

15. (Presently Amended) A method of treating endometriosis comprising administering to a patient in need thereof an effective amount of a compound of ~~any one of claims 1-14~~ claim 8, or a pharmaceutical acid addition salt thereof.

16. (Presently Amended) A method of treating uterine leiomyoma comprising administering to a patient in need thereof an effective amount of a compound of ~~any one of claims 1-14~~ claim 8, or a pharmaceutical acid addition salt thereof.

17. Cancelled

18. (Original) A compound of formula II:



II;

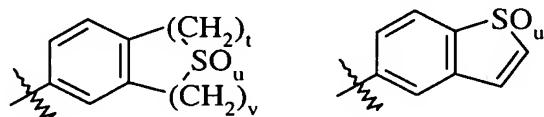
wherein:

m, q, r and u are independently 0, 1 or 2;

n is 0 or 1;

R⁰ is independently at each occurrence OH, CF₃, halo, C₁-C₆ alkyl or C₁-C₆ alkoxy;

R¹ and R^{1'} are independently C₁-C₆ alkyl, C₁-C₆ alkoxy, NR³R^{3a}, CF₃ or CH₂CF₃; or when n and q are 0, the -SO_uR¹ moiety may combine with the phenyl ring to which it is attached to form a moiety of formula (c) or (d):



(c)

(d);

wherein t and v are 0, 1 or 2 provided that the sum of t + v must be 2;

R^2 is $C_1\text{-}C_6$ alkyl; $C_1\text{-}C_6$ alkoxy; NR^4R^4 ; phenoxy; or phenyl optionally substituted with halo;

R^3 is $C_1\text{-}C_6$ alkyl or phenyl;

R^{3a} and R^4 are independently at each occurrence H, $C_1\text{-}C_6$ alkyl or phenyl;

R^6 is H, $C_1\text{-}C_6$ alkyl, benzyl or COR^2 ;

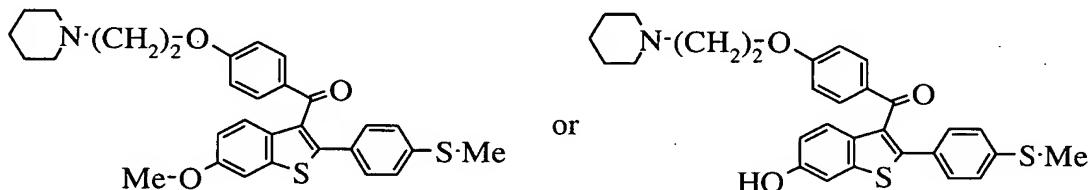
R^7 is H, $C_1\text{-}C_6$ alkyl or $CO_2(C_1\text{-}C_6$ alkyl);

R^8 is H or methyl provided that if r is 1 or 2, then R^8 must be H and that if r is 0, then R^8 must be methyl;

X is O, CH_2 or CO;

X^2 is O or NR^7 ;

Y is S, CH_2CH_2 or $CH=CH$; or a pharmaceutical acid addition salt thereof; provided that u can only be 2 when R^6 is $C_1\text{-}C_6$ alkyl or benzyl; or an acid addition salt thereof; and further provided that the compound of formula II is not:



19. (Original) The compound of claim 18, or an acid addition salt thereof, wherein r is 1 or 2; and

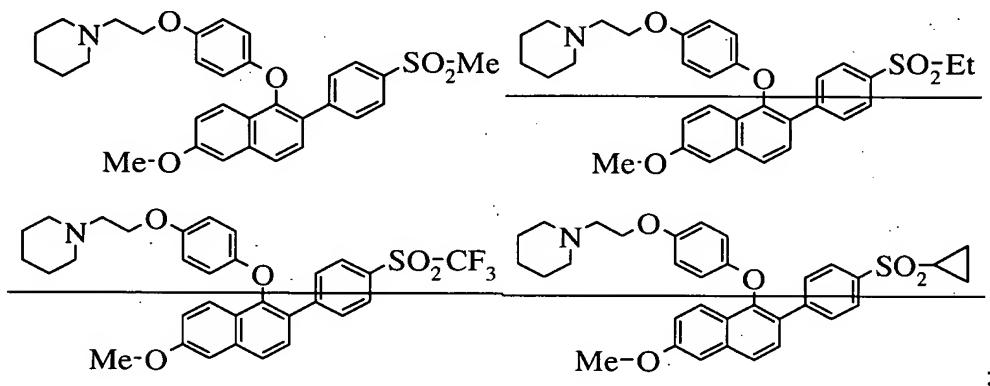
- a) if n is 0 and the SO_uR^1 moiety and R^0 combine with the phenyl ring to which they are both attached to form a moiety of formula (c) or (d), then u is 2; and
- b) if n is 1, then m and u are both 0, are both 1 or are both 2.

20. (Presently amended) The compound of ~~claim 18 or~~ 19 wherein the $-SO_uR^1$ moiety does not combine with the phenyl ring to which it is attached to form a moiety of formula (c) or (d) and is at the para-position of said phenyl ring to which it is attached; n is 0; q is 0 or 1; R^0 is OH, CF_3 , fluoro, chloro, methyl or ethyl; R^1 is methyl, ethyl, n-propyl,

isopropyl, cyclopropyl, n-butyl, isobutyl, sec-butyl, t-butyl, cyclobutyl or CF₃; R² is C₁-C₆ alkyl or phenyl; X and X¹ are O; and Y is S or CH=CH; or an acid addition salt thereof.

21. (Presently amended) The compound of any one of claims 18-20 claim 20 wherein q is 0; R¹ is methyl, ethyl, cyclopropyl or CF₃; and Y is CH=CH; or an acid addition salt thereof.

22. (Presently amended) The compound of any one of claims 18-21 selected from the group consisting of claim 20 which is:

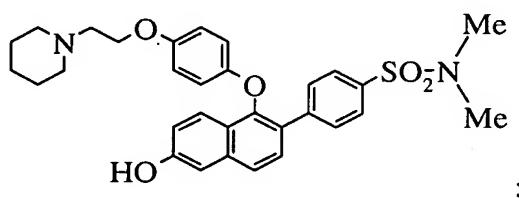


or an acid addition salt thereof.

23. (New) The method of claim 15 wherein the compound is the hydrochloride salt.

24. (New) The method of claim 16 wherein the compound is the hydrochloride salt.

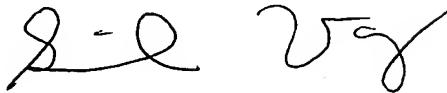
25. (New) The compound of claim 7 which is:



or a pharmaceutical acid addition salt thereof.

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Respectfully submitted,



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